10 wherein

A is an attachment linker moiety selected from the group comprising sulfur and phosphonate; MFS is a monolayer forming species comprising conductive oligomers and insulators; and AG is an electroconduit forming species.

- 2. A composition according to claim 1 wherein A is sulfur.
- 3. A composition according to claim 1 wherein said metallic surface is gold.
- 4. A composition according to claim 1 wherein said MFS is an insulator.
- 5. A composition according to claim 4 wherein said insulator comprises an alkyl group from about 7 to 20 carbons.
- 6. A composition according to claim 5 wherein said alkyl group comprises a heteroalkyl.
- 7. A composition according to claim 5 wherein said alkyl group comprises a substituted alkyl.
- 8. A composition according to claim 1 wherein said AG comprises an alkyl group from about 1 to 6 carbons.
- 9. A composition according to claim 1 or 8 wherein said AG is branched, having the formula:

wherein

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R<sub>3</sub> through R<sub>5</sub> are independently selected from the group consisting of hydrogen, alkyl, aryl, alcohol, amine, amido, nitro, ether, ester, ketone, imino, aldehyde, alkoxy, carbonyl, halogen, sulfur containing moiety and phosphorus containing moiety;

- 5 10. A composition according to claim 9 wherein said AG is attached to said attachment linker via a (CH<sub>2</sub>)<sub>n</sub> group, wherein n is an integer from 0 to 4.
  - 11. A composition according to claim 9 wherein said AG is attached directly to said attachment linker.
- 10 12. A method of modifying a metallic surface comprising contacting the metallic surface with an asymmetric monolayer forming species having the formula:



wherein

A is an attachment linker moiety;

MFS is a monlayer forming species; and

AG is an electroconduit forming species.

13. A method according to claim 12 further comprising contacting said metallic surface with a biological species having the formula:

A-MFS-capture binding ligand

wherein

A is an attachment linker; and

MFS is a monolayer forming species.

- 14. A method according to claim 13 wherein said capture binding ligand is a nucleic acid.
- 15. A method according to claim 13 wherein said capture binding ligand is a n protein.
- 35 16. A method according to claim 12 wherein A is sulfur.
  - 17. A method according to claim 12 wherein said metallic surface is gold.

20. A method according to claim 19 wherein said alkyl group comprises a heteroalkyl.

21. A method according to claim 19 wherein said alkyl group comprises a substituted alkyl.

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22. A method according to claim 12 wherein said AG comprises an alkyl group from about 1 to 6 carbons.

23. A method according to claim 12 or 22 wherein said AG is branched, having the formula:

wherein

R<sub>3</sub> through R<sub>5</sub> are independently selected from the group consisting of hydrogen, alkyl, aryl, alcohol, amine, amido, nitro, ether, ester, ketone, imino, aldehyde, alkoxy, carbonyl, halogen, sulfur containing moiety and phosphorus containing moiety;

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24. A method according to claim 23 wherein said AG is attached to said attachment linker via a  $(CH_2)_n$ group, wherein n is an integer from 0 to 4.

25. A method according to claim 23 wherein said AG is attached directly to said attachment linker.

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26. A method of detecting a target analyte in a sample comprising:

a) binding said target analyte to a metallic surface comprising

i) an asymmetric monolayer forming species having the formula:



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ii) a species having the formula A-MFS-capture binding ligand; and wherein

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A is an attachment linker moiety;

MFS is a monolayer forming species; and

AG is an electroconduit forming species; and

- b) binding a solution binding ligand to said target analyte, wherein said solution binding ligand comprises a first portion that will bind to said target analyte and a recruitment linker comprising a first portion comprising at least one ETM; and
- c) detecting the presence of said ETM as an indication of the presence of the target analyte.
- 27. A method according to claim 26 wherein said recruitment linker is directly attached to said target analyte.
- 28. A method according to claim 26 wherein said recruitment linker is indirectly attached to said target analyte.
- 29. A method according to claim 26 wherein said ETM is a transition metal complex.
- 30. A method according to claim 26 wherein said ETM is metallocene.
- 31. A method according to claim 26 wherein said ETM is ferrocene.
- 32. A method according to claim 26 wherein said ETM is an organic electron transfer moiety.
- 33. A method according to claim 26 wherein said capture binding ligand is a nucleic acid.
- 34. A method according to claim 26 wherein said capture binding ligand is a protein.
  - 35. A method according to claim 26 wherein A is sulfur.
  - 36. A method according to claim 26 wherein said metallic surface is gold.
  - 37. A method according to claim 26 wherein said MFS is an insulator.
  - 38. A method according to claim 37 wherein said insulator comprises an alkyl group from about 7 to 20 carbons.
  - 39. A method according to claim 38 wherein said alkyl group comprises a heteroalkyl.

- 40. A method according to claim 38 wherein said alkyl group comprises a substituted alkyl.
- 41. A method according to claim 26 wherein said AG comprises an alkyl group from about 1 to 6 carbons.
- 42. A method according to claim 26 wherein said AG is branched, having the formula:

## wherein

 $R_3$  through  $R_5$  is selected from the group consisting of hydrogen, alkyl, aryl, alcohol, amine, amido, nitro, ether, ester, ketone, imino, aldehyde, alkoxy, carbonyl, halogen, sulfur containing moiety and phosphorus containing moiety;

- 43. A method according to claim 26 wherein said AG is attached to said attachment linker via a  $(CH_2)_n$  group, wherein n is an integer from 0 to 4.
- 44. A method according to claim 26 wherein said AG is attached directly to said attachment linker.